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```
10-MAY-2000
                                                   BCT
                        33801 bp
LOCUS
            Streptomyces coelicolor cosmid C77.
DEFINITION
ACCESSION
            AL136503
VERSION
            AL136503.1 GI:6714747
            adenosine deaminase; carbohydrate kinase; dehydratase;
KEYWORDS
            dihydrodipicolinate synthase; DNA-binding protein; DnaJ protein;
            dnaJ2; Era-like GTP-binding protein; GTP-binding protein;
            heat-inducible transcriptional repressor; Hit-family protein; hrcA;
            hydrolase; IclR-family transcriptional regulator; lepA;
            lipoprotein; long-chain fatty-acid CoA ligase; oxidoreductase;
            oxygen-independent coproporphyrinogen III oxidase; protease;
            transmembrane efflux protein; transmembrane transport protein.
            Streptomyces coelicolor A3(2).
SOURCE
            Streptomyces coelicolor A3(2)
  ORGANISM
            Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
            Actinomycetales; Streptomycineae; Streptomycetaceae; Streptomyces.
               (bases 1 to 33801)
REFERENCE
            Redenbach, M., Kieser, H.M., Denapaite, D., Eichner, A., Cullum, J.,
  AUTHORS
            Kinashi, H. and Hopwood, D.A.
            A set of ordered cosmids and a detailed genetic and physical map
  TITLE
            for the 8 Mb Streptomyces coelicolor A3(2) chromosome
            Mol. Microbiol. 21 (1), 77-96 (1996)
  JOURNAL
  MEDLINE
            97000351
               (bases 1 to 33801)
REFERENCE
           Oliver, K. and Harris, D.
  AUTHORS
  JOURNAL
           Unpublished
               (bases 1 to 33801)
REFERENCE
           Thomson, N.R., Parkhill, J., Barrell, B.G. and Rajandream, M.A.
  AUTHORS
  TITLE
           Direct Submission
            Submitted (17-JAN-2000) Streptomyces coelicolor sequencing project,
  JOURNAL
           Sanger Centre, Wellcome Trust Genome Campus, Hinxton, Cambridge
           CB10 1SA E-mail: barrell@sanger.ac.uk Cosmids supplied by Prof.
            David A. Hopwood, [3] John Innes Centre, Norwich Research Park,
           Colney, Norwich, Norfolk NR4 7UH, UK
alignment scores:
           Quality:
                                        Length:
                                                   363
                     397.00
                                          Gaps:
                     1.829
              Ratio:
                              Percent Identity: 33.609
Percent Similarity:
                     59.780
alignment block:
US-09-477-962-115 \times SCC77/rev
Align seg 1/1 to reverse of: SCC77 from: 1 to: 33801
      11 IleProAlaIleArgGluAlaLeuGlyAspGluLysAspProArgLeuAl 27
                       :::||||||
         :::|||||
                                            111
   26678 CTGCCCGCG.....TCCGCGCTCGCCGGGGCCGCCGACCGCCCCTCGG 26635
      27 aLeuTyrValHisValProPheCysSerSerLysCysHisPheCysAspT 44
         6634 GTTCT/AGGTGGAGGTGCGTACTGCGCGACCCGCTGGGGGTACTGCGACT 26585
        rpValThrAspIleProValAlaArgLeuArgGlyAspSer
```

26584 TCAACACC. TACACCCCCACCGAACTGCGCGGCACCGGCGGGGGTCCTC 26538

 $\mathbf{H}$ 

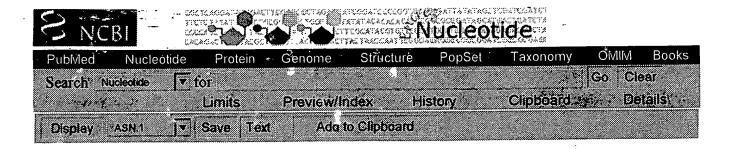
:::

A Marie Tools

	ArgGluArgSerProTyrValThrAlaLeuCysAspGlnIleArgPheTy :::       ::: ::	
26537	GCCTCCCGCGACAACTACGCCGACACCCTCGTCGACGAGGTCCGC	26493
74	rGlyProGlnLeuThrArgLeuGlyTyrArgProG	86
26492		26453
86	<pre>luValMetTyrTrpGlyGlyGlyThrProThrArgLeuThrGlyAspGlu ::::::::::::::::::::::::::::::::::::</pre>	102
26452	GCACGGTCTTCGTCGGCGGCGGTACGCCCACCCTGCTGGCCGCCGGCGAC	26403
103	MetThrAlaValHisGlnAlaLeuAspAspAlaPheAspLeuThrGlyLe :::::                :::	119
26402	CTGGTGCGGATGCTGGGCGCGATCCGCGACGAGTTCGGCCTGGCACCGGA	26353
119	uArgGlnTrpSerValGluSerThrProAsnAspLeuAspProAlaThrL ::: :::::   :::::   ::::	136
26352	CGCGGAGATCACCACGGAGGCCAACCCGGAGTCCGTCGACCCGGCGTATC	26303
136	euAspThrLeuArgGlyLeuGlyValThrArgValSerValGlyValGln	152
26302	TCGCCACCCTCCGCGGGGGGGGCTTCAACCGGATCTCCTTCGGCATGCAG	26253
153	SerLeuAsnProTyrGlnLeuArgLysAlaGlyArgAlaHisSerArgGl	169
26252	AGCGCCAAGCACGCCCTGAAGATCCTGGACCGCACCCACACCCCGGG	26203
169	uGlnAlaLeuAlaAlaValProLeuLeuArgArgAlaGlyIleAspGluP :::    :::               :::   :::	186
26202	ACGCCCGAGGCCTGTGTCGCCGAGGCCCGCGCGGCCGGCTTCGACCACG	26153
186	heAsnValAspLeuIleAlaGlyPheProGlyGluAlaValGluSerPhe	202
26152	TCAACCTCGACCTGATCTACGGCACCCCCGGCGAGTCCGACGACGACTGG	26103
203	GluGluThrLeuArgThrValLeuAlaLeuAspProProHisValSerVa ::: :::   :::::  :::     :	219
26102	CGGGCCTCCCTGGACGCCGCGCTCGGCGCCCGACCACGTCTCGGC	26053
219	<pre>lTyrProTyrArgAlaThrProLysThrValMetAlaMetGlnLeuAspA :    :::     ::::! </pre>	236
26052	GTACGCCTGATCGTCGAGGAGGGCACCCAGCTCGCCCGCC	26003
236	rgGluPheValGluAlaArgAsnArgAspGlyMetIleAspAlaTyrGlu	252
26002	GCGGCGAGGTCCCGATGACCGACGACGACGTGCACGCCGACCGGTACCTG	25953
253	ArgAlaMetAlaAlaLeuGlyAlaAlaGlyTyrHisGluTyrCysHisGl	269
1.1	ATCGCCGAGGAGGCACTGTCCGCGGCCGGTTACGACTGGTACGAGGTGTC	
269	YTyrTrpValArgAspAlaArgHisGluAspGlnAspG	282
	CAACTGGGCCACCTCCGACGCGGGGCGCTGCCTGCAC	25866
A STATE OF THE STA	respectively and the control of the	

AND THE PROPERTY OF THE PROPER

282	lyAsnTyrLysTyrAspLeuAlaGlyAspLysIleGlyPheGlySerGly	298
25865	AACGAGCTGTACTGGCGGGGCGCCGACTGGTGGGGCGCGGGACCGGGC	25818
299	${\tt AlaGluSerIleIleGlyHisHisLeuLeuTrpAsnGluAsnSe}$	313
	)	
25817	GCGCACTCCCACGTGGGGGGCGTGCGGTGGTGGAACGTGAAGCACCCGGG	25768
313	rAlaTyrAlaArgTyrLeuLeuAlaProArgGluPheSerAlaAlaHisA	330
	:	05710
25767	GGCGTACGCGGGGGCGCTGGCGGGGGGGCGCGGGGCGCGGGGCGCG	25/18
		246
330-	rgPheThrThrAlaGluProAspArgLeuThrAlaProValGlyGlyAla	346
05515	:::::             ::: :::	25671
25/1/	AGATCCTCACGGACGAGGACCGGCGCGTGGAGCGCATCCTGCTGGAG	236/1
247	Towns to the control of the late of the la	
34/	LeuMetThrArgGluGlyValValPheAlaArgPheArg 359	
05.670	:::::: :::	
256/0	CTGCGCCTGCGGGAGGGCGTCCCGCTGTCGCTGCTGCGG 25632	



I: AL136503. Streptomyces coel...[gi:6714747] Related Sequences, Protein, PubMed, Taxonomy

LOCUS SCC77 33801 bp DNA linear BCT 10-MAY-2000

DEFINITION Streptomyces coelicolor cosmid C77.

ACCESSION AL136503

VERSION AL136503.1 GI:6714747

KEYWORDS adenosine deaminase; carbohydrate kinase; dehydratase;

dihydrodipicolinate synthase; DNA-binding protein; DnaJ protein;

dnaJ2; Era-like GTP-binding protein; GTP-binding protein;

heat-inducible transcriptional repressor; Hit-family protein; hrcA;

hydrolase; IclR-family transcriptional regulator; lepA;

lipoprotein; long-chain fatty-acid CoA ligase; oxidoreductase; oxygen-independent coproporphyrinogen III oxidase; protease; transmembrane efflux protein; transmembrane transport protein.

SOURCE Streptomyces coelicolor A3(2).
ORGANISM Streptomyces coelicolor A3(2)

Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;

Actinomycetales; Streptomycineae; Streptomycetaceae; Streptomyces.

REFERENCE 1 (bases 1 to 33801)

AUTHORS Redenbach, M., Kieser, H.M., Denapaite, D., Eichner, A., Cullum, J.,

Kinashi, H. and Hopwood, D.A.

TITLE A set of ordered cosmids and a detailed genetic and physical map

for the 8 Mb Streptomyces coelicolor A3(2) chromosome

JOURNAL Mol. Microbiol. 21 (1), 77-96 (1996)

MEDLINE 97000351

REFERENCE 2 (bases 1 to 33801)

AUTHORS Oliver, K. and Harris, D.

JOURNAL Unpublished

REFERENCE 3 (bases 1 to 33801)

AUTHORS Thomson, N.R., Parkhill, J., Barrell, B.G. and Rajandream, M.A.

TITLE Direct Submission

JOURNAL Submitted (17-JAN-2000) Streptomyces coelicolor sequencing project,

Sanger Centre, Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SA E-mail: barrell@sanger.ac.uk Cosmids supplied by Prof. David A. Hopwood, [3] John Innes Centre, Norwich Research Park,

Colney, Norwich, Norfolk NR4 7UH, UK

COMMENT Notes:

Streptomyces coelicolor sequencing at The Sanger Centre is funded

by the BBSRC and Beowulf Genomics

Details of S. coelicolor sequencing at the Sanger Centre are

available on the World Wide Web.

(URL; http://www.sanger.ac.uk/Projects/S\_coelicolor/)

CDS are numbered using the following system eg SC7B7.01c. SC (S. coelicolor), 7B7 (cosmid name), .01 (first CDS), c (complementary strand).

The more significant matches with motifs in the PROSITE database are also included but some of these may be fortuitous.

The length in codons is given for each CDS.

Usually the highest scoring match found by fasta -o is given for CDS which show significant similarity to other CDS in the database. The position of possible ribosome binding site sequences are given

where these have been used to deduce the initiation codon.

Gene prediction is based on positional base preference in codons



# Protein

PubMed - I	Nucleotide	Protein	Genome	Structure	PopSet	Taxonomy	OMIMO	Books
Search Nucle	otide 🔻 t	or					Go Cle	ar
		Limits	Preview/In	dex His	story	Clipboard	De	tails
Display de	fault 🔻	Save   Text	Add t	Clipboard				

**1:** CAB66237. putative oxygen-i...[gi:6714773]

Nucleotide, Related Sequences, PubMed, Taxonomy, BLink, LinkOut

BCT 10-MAY-2000 435 aa linear LOCUS CAB66237 DEFINITION putative oxygen-independent coproporphyrinogen III oxidase. [Streptomyces coelicolor A3(2)]. ACCESSION CAB66237 PID g6714773 VERSION CAB66237.1 GI:6714773 **DBSOURCE** embl locus SCC77, accession AL136503.1 KEYWORDS Streptomyces coelicolor A3(2). SOURCE Streptomyces coelicolor A3(2) ORGANISM Bacteria; Firmicutes; Actinobacteria; Actinobacteridae; Actinomycetales; Streptomycineae; Streptomycetaceae; Streptomyces. REFERENCE 1 (residues 1 to 435) Redenbach, M., Kieser, H.M., Denapaite, D., Eichner, A., Cullum, J., AUTHORS Kinashi, H. and Hopwood, D.A. TITLE A set of ordered cosmids and a detailed genetic and physical map for the 8 Mb Streptomyces coelicolor A3(2) chromosome Mol. Microbiol. 21 (1), 77-96 (1996) JOURNAL. 97000351 MEDLINE REFERENCE (residues 1 to 435) AUTHORS Oliver, K. and Harris, D. JOURNAL Unpublished REFERENCE (residues 1 to 435) Thomson, N.R., Parkhill, J., Barrell, B.G. and Rajandream, M.A. **AUTHORS** TITLE Direct Submission Submitted (17-JAN-2000) Streptomyces coelicolor sequencing project, JOURNAL Sanger Centre, Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SA E-mail: barrell@sanger.ac.uk Cosmids supplied by Prof. David A. Hopwood, [3] John Innes Centre, Norwich Research Park, Colney, Norwich, Norfolk NR4 7UH, UK COMMENT Notes: Streptomyces coelicolor sequencing at The Sanger Centre is funded by the BBSRC and Beowulf Genomics Details of S. coelicolor sequencing at the Sanger Centre are available on the World Wide Web. (URL; http://www.sanger.ac.uk/Projects/S\_coelicolor/) CDS are numbered using the following system eg SC7B7.01c. SC (S. coelicolor), 7B7 (cosmid name), .01 (first CDS), c (complementary strand). The more significant matches with motifs in the PROSITE database are also included but some of these may be fortuitous. The length in codons is given for each CDS. Usually, the highest scoring match found by fasta -o is given for-CDS which show significant similarity to other CDS in the database. The position of possible ribosome binding site sequences are given where these have been used to deduce the initiation codon. Gene prediction is based on positional base preference in codons using a specially developed Hidden Markov Model (Krogh et al., Nucleic Acids Research, 22(22):4768-4778(1994)) and the FramePlot program of Bibb et al., Gene 30:157-66(1984) as implemented at

以我们的一个人,我们就是一个人的一个人的一个人,我们就是一个人的一个人的,我们就是一个人的一个人的一个人的一个人的一个人的一个人的一个人的一个人的一个人的一个人

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http://www.nih.go.jp/
            jun/cgi-bin/frameplot.pl. CAUTION: We may not have predicted the
            correct initiation codon. Where possible we choose an initiation
            codon (atg, gtg, ttg or (att)) which is preceded by an upstream
            ribosome binding site sequence (optimally 5-13bp before the
            initiation codon). If this cannot be identified we choose the most
            upstream initiation codon.
            IMPORTANT: This sequence MAY NOT be the entire insert of the
            sequenced clone. It may be shorter because we only sequence
            overlapping sections once, or longer, because we arrange for a
            small overlap between neighbouring submissions.
            Cosmid C77 Lies between and overlaps with cosmids C117 and C123 on
            the AseI-C genomic restriction fragment.
FEATURES
                     Location/Qualifiers
                     1..435
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                     /organism="Streptomyces coelicolor A3(2)"
                     /strain="A3(2)"
                     /db xref="taxon:100226"
                     /clone="cosmid C77"
     Protein
                     1..435
                     /product="putative oxygen-independent coproporphyrinogen
                     III oxidase."
     CDS
                     1..435
                     /gene="SCC77.26c"
                     /coded by="complement(AL136503.1:25494..26801)"
                     /transl table=11
                     /note="SCC77.26c, possible oxygen-independent
                     coproporphyrinogen III oxidase (EC 1.-.-.), len: 435 aa.
                     Highly similar to many putative coproporphyrinogen III
                     oxidases including: Bacillus subtilis
                     SW:HEMN BACSU(EMBL:X91655) probable oxygen-independent
                     coproporphyrinogen III oxidase (366 aa), fasta scores opt:
                     490 z-score: 555.3 E(): 1.5e-23 30.8% identity in 328 aa
                     overlap and Mycobacterium tuberculosis
                     SW:HEMN MYCTU(EMBL:Z81368) probable oxygen-independent
                     coproporphyrinogen III oxidase (375 aa), fasta scores opt:
                     1358 z-score: 1530.5 E():0 56.5% identity in 382 aa
                     overlap. Contains a Prosite hit to PS00017 ATP/GTP-binding
                     site motif A (P-loop)."
ORIGIN
        1 mngrreraqg tewagdpagc gtmermpsal pdgepvpadg alpasalaga adrplgfylh
       61 vpycatrcgy cdfntytate lrgtggvlas rdnyadtlvd evrlarkvlg ddprevrtvf
      121 vgggtptlla agdlvrmlga irdefglapd aeitteanpe svdpaylatl raggfnrisf
      181 gmqsakqhvl kildrthtpg rpeacvaear aagfdhvnld liygtpgesd ddwrasldaa
     241 lgagpdhvsa yaliveegtq larrirrgev pmtdddvhad ryliaeeals aagydwyevs
     301 nwatsdagrc lhnelywrga dwwgagpgah shvggvrwwn vkhpgayaga laagkspgag
     361 reiltdedrr verillelrl regvplsllr eaglaasrra lsegllqegp yeagsavltl
      421 rgrlladavv rdlvd
11
```

Revised: October 24, 2001.

5 x US-09-477-962-1 ...

Seg 1= 58857 6p 57,583-58,854

alignment block:  $us-09-477-962-115 \times US-09-477-962-1$ Align seg 1/1 to: US-09-477-962-1 from: 1 to: 58857 1 MetSerHisAlaIleGlyProSerArgLeuIleProAlaIleArgGluAl 17 57583 ATGAGCCACGCCATCGGAPCGAGCCGGCTGATCCCCGCCATCCGCGAAGC 57632 17 aLeuGlyAspGluLysAspProArgLeuAlaLeuTyrValHisValProP 34 57633 GCTCGGGGACGAGAAGGACCCCCGGCTCGCCCTCTACGTCCACGTCCCCT 57682 34 heCysSerSerLysCysHisPheCysAspTrpValThrAspIleProVal 50 57683 TCTGCTCCTCCAAGTGCCACTTCTGCGACTGGGTCACCGACATCCCCGTC 57732 51 AlaArgLeuArgGlyAspSerArgGluArgSerProTyrValThrAlaLe 67 57733 GCACGCCTGCGCGGCACAGCCGGGAACGCTCGCCCTACGTCACCGCCCT 57782 67 uCysAspGlnIleArgPheTyrGlyProGlnLeuThrArgLeuGlyTyrA 84 57783 CTGCGACCAGATCCGCTTCTACGGCCCCCAGCTCACCCGGCTCGGCTACC 57832 84 rgProGluValMetTyrTrpGlyGlyGlyThrProThrArgLeuThrGly 100 57833 GCCCCGAGGTCATGTACTGGGGCGGCGCCACCCCACCCGGCTCACCGGC 57882 101 AspGluMetThrAlaValHisGlnAlaLeuAspAspAlaPheAspLeuTh 117 57883 GACGAGATGACGGCCGTCCACCAGGCCCTCGACGACGCCTTCGACCTGAC 57932 117 rGlyLeuArgGlnTrpSerValGluSerThrProAsnAspLeuAspProA 134 57933 GGGACTCCGCCAGTGGTCGGTGGAGAGCACCCCGAACGACCTCGACCCCG 57982 134 laThrLeuAspThrLeuArgGlyLeuGlyValThrArgValSerValGly 150 57983 CCACCCTCGACACCCTGCGCGCGCCTCACCCGCGTCAGCGTCGGC 58032 151 ValGlnSerLeuAsnProTyrGlnLeuArgLysAlaGlyArgAlaHisSe 167 58033 GTCCAGTCGCTCAACCCGTACCAGCTGCGCAAGGCAGGCCGGGCCCACTC 58082 167 rArgGluGlnAlaLeuAlaAlaValProLeuLeuArgArgAlaGlyIleA 184 58083 GCGCGAACAGGCCCTGGCCGCCGTCCCCCTGTTGCGCCGCCGCCGCATCG 58132 184\_spGluPheAsnValAspLeuIleAlaGlyPheProGlyGluAlaValGlu 200 CACTITICAN GET GENECIT GAT COCCEGGET TICE GE GGG GANG GE GT GGA GA S 8 1.8 2 ....

201 SerPheGluGluThrLeuArgThrValLeuAlaLeuAspProProHisVa 217

217	lSerValTyrProTyrArgAlaThrProLysThrValMetAlaMetGlnL	234
	CTCCGTCTACCCCTACCGCGCCACCCCCAAGACGGTCATGGCCATGCAGC	58282
234	euAspArgGluPheValGluAlaArgAsnArgAspGlyMetIleAspAla	250
58283	TCGACCGCGAGTTCGTCGAGGCCCGGAACCGGGACGGCATGATCGACGCC	58332
251	TyrGluArgAlaMetAlaAlaLeuGlyAlaAlaGlyTyrHisGluTyrCy	267
58333	TATGAACGGGCCATGGCCGCCGCCGCCGGCTATCACGAGTACTG	58382
267	sHisGlyTyrTrpValArgAspAlaArgHisGluAspGlnAspGlyAsnT	284
58383	CCACGGCTACTGGGTGCGCGACGCGCCACGAGGACCAGGACGCCAACT	58432
284	<pre>yrLysTyrAspLeuAlaGlyAspLysIleGlyPheGlySerGlyAlaGlu</pre>	300
58433	ACAAGTACGACCTGGCCGGCGACAAGATCGGCTTTGGCAGCGGCGCCGAA	58482
301	SerIleIleGlyHisHisLeuLeuTrpAsnGluAsnSerAlaTyrAlaAr	317
58483	TCGATCATCGGTCACCACCTGCTCTGGAACGAGAACAGCGCCTACGCCCG	58532
317	gTyrLeuLeuAlaProArgGluPheSerAlaAlaHisArgPheThrThrA	334
58533	CTACCTGCTCGCCCCCGCGAGTTCTCCGCCGCCCACCGGTTCACCACCG	58582
334	laGluProAspArgLeuThrAlaProValGlyGlyAlaLeuMetThrArg	350
58583	CCGAACCCGACCGCCTGACCGCCCCCGTCGGCGCGCGCTGATGACCCGT	58632
351	GluGlyValValPheAlaArgPheArgArgLeuThrGlyLeuAspPheAl	3.67
58633	GAAGGCGTGGTCTTCGCCCGCTTCCGCAGACTGACCGGCCTGGACTTCGC	58682
367	aAspValArgAlaThrProTyrPheArgGlnTrpPheGluLeuLeuGluA	384
58683	GGACGTCCGCCACACCGTACTTCCGCCAGTGGTTCGAGCTCCTGGAGC	58732
384	rgCysGlyGlyArgPheValGluThrProTyrSerLeuArgLeuGluPro	400
58733	GCTGCGGCGGCCGCTTCGTCGAGACGCCGTACAGCCTCCGCCTGGAGCCG	58782
401	SerThrIleHisArgAlaTyrIleThrHisLeuAlaTyrThrMetAlaHi	417
58783	TCCACCATCCACCGCGCCTACATCACCCACCTCGCCTACACCATGGCCCA	58832
417	sGlyLeuAlaProGluArgAla 424	
58833	TGGCCTGGCCCCGAACGCGCC 58854	ing in Literatus

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And the second second

- L1 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS
- AN 2000:332950 CAPLUS
- TI Bleomycin biosynthesis in Streptomyces verticillus ATCC15003: A model for hybrid peptide and polyketide biosynthesis.
- AU Du, Liangcheng; Sanchez, Cesar; Chen, Mei; Edwards, Daniel J.; Murrell, Jeffrey M.; Shen, Ben
- CS Department of Chemistry, University of California, Davis, CA, 95616, USA
- SO Book of Abstracts, 219th ACS National Meeting, San Francisco, CA, March 26-30, 2000 (2000), ORGN-822 Publisher: American Chemical Society, Washington, D. C.
  - CODEN: 69CLAC
- DT Conference; Meeting Abstract
- LA English

#### => d abs

- L1 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS
- AB Polyketides and nonribosomal peptides are assembled in a remarkably similar fashion by polyketide synthases (PKSs) from short carboxylic acids and nonribosomal peptide synthetases (NRPSs) from amino acids, resp. Cloning and sequence anal. of the 90-kb bleomycin (BLM) biosynthesis cluster from Streptomyces verticillus ATCC15003 revealed both NRPS and PKS genes. By detg. the substrate specificity of individual NRPS and PKS modules, a linear hybrid NRPS/PKS/NRPS model is formulated for the Blm megasynthetase-templated assembly of BLM from nine amino acids and one acetate. These results set the stage for engineering novel BLM analogs by genetic manipulation of the blm biosynthesis genes, support the wisdom of combining individual NRPS and PKS modules for combinatorial biosynthesis, and lay the foundation to investigate the mol. basis for intermodular communication between NRPS and PKS and the mechanism for bithiazole biosynthesis.

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- L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS
- AN 1999:145586 CAPLUS
- TI Bleomycin biosynthesis in streptomyces verticillus ATCC15003: The search for a hybrid polyketide and peptide biosynthetic system
- AU Shen, Ben; Du, Liangcheng; Edwards, Dan; Chen, Mei; Sanchez, Cesar
- CS Department of Chemistry, University of California, Davis, CA, 95616, USA
- SO Book of Abstracts, 217th ACS National Meeting, Anaheim, Calif., March 21-25 (1999), ORGN-153 Publisher: American Chemical Society, Washington, D. C.
  - CODEN: 67GHA6
- DT Conference; Meeting Abstract
- LA English

#### => d abs

- L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS
- AB The biosynthesis of bleomycin (Blm) in Sv. ATCC15003 has been studied as a model to decipher the mechanism of how peptide synthase (PTS) and polyketide synthase (PKS) can be hybridized into a functional system to make metabolite from amino acids and short fatty acids. A 110kb gene cluster for Blm biosynthesis was cloned from Sv. ATCC15003, 75kb of which has been fully sequenced and analyzed. Among the many novel discoveries made from this study are:. (1) the first model for a hybrid PTS/PKS/PTS biosynthetic system, (2) the first example of PKS gene from actinomycetes that contains a MT domain, and (3) a novel mechanism for bithiazole biosynthesis. These results should lay the foundation for rational engineering of hybrid PTS/PKS biosynthetic systems from other peptide and polyketide biosynthetic pathways to generate structural diversity.

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